

**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of the Claims:**

1. (Currently amended) An isolated polypeptide sequence selected from the group consisting of:
  - a) a polypeptide comprising the ~~an~~ amino acid sequence of SEQ ID NO:1, and
  - b) a polypeptide that is ~~comprising a naturally occurring amino acid sequence~~ at least 95% ~~[[90%]]~~ identical to the ~~an~~ amino acid sequence of SEQ ID NO:1, said polypeptide having DNA binding activity;
  - ~~e) a biologically active fragment of a polypeptide having an amino acid sequence of SEQ ID NO:1, said polypeptide having DNA binding activity, and~~
  - ~~d) an immunogenic fragment of a polypeptide having an amino acid sequence of SEQ ID NO:1.~~
2. (Previously presented) An isolated polypeptide of claim 1, comprising the amino acid sequence of SEQ ID NO:1.
- 3.-13. (Canceled)
14. (Withdrawn) A method for detecting a target polynucleotide in a sample, said target polynucleotide having a sequence of a polynucleotide of claim 12, the method comprising:
  - a) hybridizing the sample with a probe comprising at least 20 contiguous nucleotides comprising a sequence complementary to said target polynucleotide in the sample, and which probe specifically hybridizes to said target polynucleotide, under conditions whereby a hybridization complex is formed between said probe and said target polynucleotide or fragments thereof, and
  - b) detecting the presence or absence of said hybridization complex, and, optionally, if present, the amount thereof.

15. (Withdrawn) A method of claim 14, wherein the probe comprises at least 60 contiguous nucleotides.
16. (Withdrawn) A method for detecting a target polynucleotide in a sample, said target polynucleotide having a sequence of a polynucleotide of claim 12, the method comprising:
- a) amplifying said target polynucleotide or fragment thereof using polymerase chain reaction amplification, and
  - b) detecting the presence or absence of said amplified target polynucleotide or fragment thereof, and, optionally, if present, the amount thereof.
17. (Original) A composition comprising a polypeptide of claim 1 and a pharmaceutically acceptable excipient.
- 18 (Previously presented) A composition of claim 17, wherein the polypeptide comprises an amino acid sequence of SEQ ID NO:1.
19. (Withdrawn) A method for treating a disease or condition associated with decreased expression of functional PRAEP, comprising administering to a patient in need of such treatment the composition of claim 17.
20. (Withdrawn) A method for screening a compound for effectiveness as an agonist of a polypeptide of claim 1, the method comprising:
- a) exposing a sample comprising a polypeptide of claim 1 to a compound, and
  - b) detecting agonist activity in the sample.
21. (Withdrawn) A composition comprising an agonist compound identified by a method of claim 20 and a pharmaceutically acceptable excipient.
22. (Withdrawn) A method for treating a disease or condition associated with decreased expression of functional PRAEP, comprising administering to a patient in need of such treatment a composition of claim 21.

23. (Withdrawn) A method for screening a compound for effectiveness as an antagonist of a polypeptide of claim 1, the method comprising:

- a) exposing a sample comprising a polypeptide of claim 1 to a compound, and
- b) detecting antagonist activity in the sample.

24. (Withdrawn) A composition comprising an antagonist compound identified by a method of claim 23 and a pharmaceutically acceptable excipient.

25. (Withdrawn) A method for treating a disease or condition associated with overexpression of functional PRAEP, comprising administering to a patient in need of such treatment a composition of claim 24.

26. (Withdrawn) A method of screening for a compound that specifically binds to the polypeptide of claim 1, said method comprising:

- a) combining the polypeptide of claim 1 with at least one test compound under suitable conditions, and
- b) detecting binding of the polypeptide of claim 1 to the test compound, thereby identifying a compound that specifically binds to the polypeptide of claim 1.

27. (Withdrawn) A method of screening for a compound that modulates the activity of the polypeptide of claim 1, said method comprising:

- a) combining the polypeptide of claim 1 with at least one test compound under conditions permissive for the activity of the polypeptide of claim 1,
- b) assessing the activity of the polypeptide of claim 1 in the presence of the test compound, and
- c) comparing the activity of the polypeptide of claim 1 in the presence of the test compound with the activity of the polypeptide of claim 1 in the absence of the test compound,

wherein a change in the activity of the polypeptide of claim 1 in the presence of the test compound is indicative of a compound that modulates the activity of the polypeptide of claim 1.

28. (Withdrawn) A method for screening a compound for effectiveness in altering expression of a target polynucleotide, wherein said target polynucleotide comprises a polynucleotide sequence of claim 5, the method comprising:

- a) exposing a sample comprising the target polynucleotide to a compound, under conditions suitable for the expression of the target polynucleotide,
- b) detecting altered expression of the target polynucleotide, and
- c) comparing the expression of the target polynucleotide in the presence of varying amounts of the compound and in the absence of the compound.

29. (Withdrawn) A method for assessing toxicity of a test compound, said method comprising:

- a) treating a biological sample containing nucleic acids with the test compound;
- b) hybridizing the nucleic acids of the treated biological sample with a probe comprising at least 20 contiguous nucleotides of a polynucleotide of claim 12 under conditions whereby a specific hybridization complex is formed between said probe and a target polynucleotide in the biological sample, said target polynucleotide comprising a polynucleotide sequence of a polynucleotide of claim 12 or fragment thereof;
- c) quantifying the amount of hybridization complex; and
- d) comparing the amount of hybridization complex in the treated biological sample with the amount of hybridization complex in an untreated biological sample, wherein a difference in the amount of hybridization complex in the treated biological sample is indicative of toxicity of the test compound.

30.-45. (Canceled)

46. (Original) A polypeptide of claim 1, comprising the amino acid sequence of SEQ ID NO:1.

47. (Withdrawn) A polynucleotide of claim 12, comprising the polynucleotide sequence of SEQ ID NO:2.